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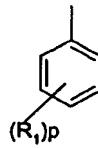
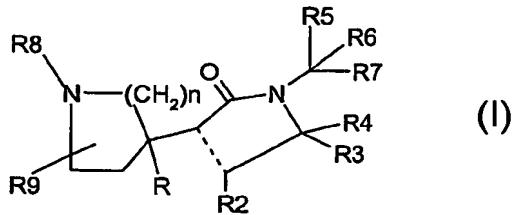
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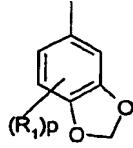
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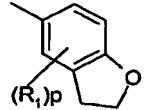
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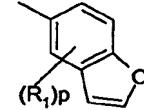
(i)



(ii)



(iii)



(iv)

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(57) Abstract: The present invention relates to novel compounds of formula (I) wherein ---- represents a single or a double bond; R represents a radical selected from formulae i), ii), iii) and iv) in which R₁ is halogen, cyano, C₁₋₄ alkyl, C₁₋₄ alkoxy, trifluoromethyl or trifluoromethoxy and p is zero or an integer from 1 to 3; R₂ represents hydrogen or C₁₋₄ alkyl; R₃ represents hydrogen, hydroxy or C₁₋₄ alkyl; R₄ represents hydrogen or R₄ together with R₃ represents =O or =CH₂; R₅ represents phenyl, naphthyl, a 9 to 10 membered fused bicyclic heterocyclic group or a 5 or 6 membered heteroaryl group, wherein said groups are optionally substituted by 1 to 3 groups independently selected from trifluoromethyl, C₁₋₄ alkyl, hydroxy, cyano, C₁₋₄ alkoxy, trifluoromethoxy, halogen or S(O)_qC₁₋₄ alkyl; R₆ and R₇ independently represent hydrogen, cyano, C₁₋₄ alkyl; R₈ is (CH₂)_rR₁₀; R₉ represents hydrogen, halogen, C₃₋₇ cycloalkyl, hydroxy, nitro, cyano or C₁₋₄ alkyl optionally substituted by one or two groups selected from halogen, cyano, hydroxy or C₁₋₄ alkoxy; R₁₀ represents hydrogen or C₃₋₇ cycloalkyl; n represents 1 or 2; q is 0, 1 or 2; r is 0 or an integer from 1 to 4; or a pharmaceutically acceptable salt or a solvate thereof, process for their preparation and their use in the treatment of conditions mediated by tachykinins and/or by selective inhibition of the serotonin reuptake transporter protein.



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